

MUCUS RELIEF DM- dextromethorphan hydrobromide guaifenesin liquid

The Kroger Co.

Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg

Guaifenesin 400 mg

Purpose

Cough suppressant

Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive
- temporarily relieves
 - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
 - the intensity of coughing
 - the impulse to cough to help you get to sleep

Warnings

Do not use

if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough accompanied by too much phlegm (mucus)

When using this product,

do not use more than directed.

Stop use and ask a doctor if

cough lasts more than 7 days, comes back, or occurs with fever, rash or persistent headache. These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other device
- keep dosing cup with product
- mL = milliliter
- dose as follows or as directed by a doctor
- adults and children 12 years of age and older: 20 mL every 4 hours
- children under 12 years of age: do not use

Other information

- **each 20 mL contains:** sodium 20 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

citric acid, disodium EDTA, FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

Questions or comments?

Call 1-800-632-6900

Principal Display Panel

COMPARE TO the active ingredients of MUCINEX® FAST-MAX® DM MAX

for ages 12+

maximum Strength

FAST

Mucus Relief

DM Max

multi-Symptom

Dextromethorphan HBr 20 mg

Cough Suppressant

Guaifenesin 400 mg

Expectorant

Relieves

Chest Congestion

Controls Cough

Thin & Loosens Mucus

4 Hour Dosing

*Mucinex® Fast-Max® DM Max is a registered trademark of Reckitt Benckiser, Slough, UK. Reckitt Benckiser is not affiliated with The Kroger Co. or this product

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS
BROKEN OR MISSING.**

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CINCINNATI, OHIO 45202

www.kroger.com

Package Label

COMPARE TO the active ingredients of
MUCINEX® FAST-MAX® DM MAX

*See back panel

NDC 30142-508-06



for ages 12+



Maximum Strength

FAST Mucus Relief DM Max

Multi-Symptom

Dextromethorphan HBr
Cough Suppressant

Guaifenesin
Expectorant

Relieves
Chest Congestion

Controls Cough

Thins & Loosens Mucus

4 Hour Dosing

6 FL OZ (177 mL)

PLD-A409A LB008052

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY
SEAL AROUND OR UNDER CAP IS BROKEN OR MISSING.**

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PARENTS:
Learn about teen medicine abuse
www.StopMedicineAbuse.org

PLD-A409A LB008053



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CINCINNATI, OHIO 45202
QUALITY GUARANTEE
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PEEL CORNER FOR DRUG FACTS

Drug Facts

Active ingredients Purposes (in each 20 mL)

Dextromethorphan HBr 20 mg.....
.....Cough suppressant
Guaifenesin 400 mg.....Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves
 - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
 - the intensity of coughing
 - the impulse to cough to help you get to sleep

Drug Facts (continued)

Warnings

Do not use if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)

When using this product, do not use more than directed.

Stop use and ask a doctor if cough lasts more than 7 days, comes back, or occurs with fever, rash, or headache that lasts. These could be signs of a serious condition.

Drug Facts (continued)

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- do not take more than 6 doses in any 24-hour period
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- mL = milliliter
- dose as follows or as directed by a doctor
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Other information

- each 20 mL contains: sodium 20 mg
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Drug Facts (continued)

Inactive ingredients

citric acid, disodium EDTA, FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

Questions or comments?

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PEEL CORNER FOR MORE DRUG FACTS

KROGER Maximum Strength Mucus Relief DM

MUCUS RELIEF DM

dextromethorphan hydrobromide guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-508
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KOOR)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-508-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/01/2020	

Labeler - The Kroger Co. (006999528)

